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FILE COVERS 1971 TO PATENT PUBLICATION DATE: 27 May 2003 (20030527/PD)
FILE LAST UPDATED: 27 May 2003 (20030527/ED)
HIGHEST GRANTED PATENT NUMBER: US6571393
HIGHEST APPLICATION PUBLICATION NUMBER: US2003097700
CA INDEXING IS CURRENT THROUGH 27 May 2003 (20030527/UPCA)
ISSUE CLASS FIELDS (/INCL) CURRENT THROUGH: 27 May 2003 (20030527/PD)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Feb 2003
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Feb 2003

USPAT2 is now available. USPATFULL contains full text of the >>> <<< original, i.e., the earliest published granted patents or <<< >>> <<< applications. USPAT2 contains full text of the latest US >>> <<< publications, starting in 2001, for the inventions covered in >>> <<< USPATFULL. A USPATFULL record contains not only the original >>> <<< >>> published document but also a list of any subsequent >>> publications. The publication number, patent kind code, and <<< >>> publication date for all the US publications for an invention <<< >>> are displayed in the PI (Patent Information) field of USPATFULL <<< >>> records and may be searched in standard search fields, e.g., /PN, <<< <<< >>> /PK, etc. USPATFULL and USPAT2 can be accessed and searched together <<< through the new cluster USPATALL. Type FILE USPATALL to <<< >>> >>> enter this cluster. <<< <<< >>> Use USPATALL when searching terms such as patent assignees, <<< classifications, or claims, that may potentially change from <<< >>> the earliest to the latest publication. <<<

This file contains CAS Registry Numbers for easy and accurate substance identification.

L6 277 CIGLITAZONE AND 5 MG

```
=> s ciglitazone (1P) 5 mg
          414 CIGLITAZONE
      3340772 5
       306382 MG
        88222 5 MG
                 (5 (W) MG)
L7
           26 CIGLITAZONE (1P) 5 MG
=> d 17 and pd<1999
'AND' IS NOT A VALID FORMAT FOR FILE 'USPATFULL'
'PD<1999' IS NOT A VALID FORMAT FOR FILE 'USPATFULL'
The following are valid formats:
The default display format is STD.
ABS ----- AB
ALL ----- AN, TI, IN, INA, PA, PAA, PAT, PI, AI, PTERM, DCD,
            RLI, PRAI, DT, FS, REP, REN, EXNAM, LREP, CLMN, ECL,
            DRWN, AB, GOVI, PARN, SUMM, DRWD, DETD, CLM, INCL,
            INCLM, INCLS, NCL, NCLM, NCLS, IC, ICM, ICS,
            EXF, ARTU
ALLG ----- ALL plus PAGE.DRAW
BIB ----- AN, TI, IN, INA, PA, PAA, PAT, PI, AI, PTERM, DCD, RLI,
            PRAI, DT, FS, EXNAM, LREP, CLMN, ECL, DRWN, LN.CNT
BIB.EX ---- BIB for original and latest publication
BIBG ----- BIB plus PAGE.DRAW
BROWSE ---- See "HELP BROWSE" or "HELP DISPLAY BROWSE". BROWSE must
            entered on the same line as DISPLAY, e.g., D BROWSE.
CAS ----- OS, CC, SX, ST, IT
CBIB ----- AN, TI, IN, INA, PA, PAA, PAT, PI, AI, PRAI, DT, FS
DALL ---- ALL, delimited for post-processing
FP ----- PI, TI, IN, INA, PA, PAA, PAT, PTERM, DCD, AI, RLI,
            PRAI, IC, ICM, ICS, INCL, INCLM, INCLS, NCL,
            NCLM, NCLS, EXF, REP, REN, ARTU, EXNAM, LREP,
            CLMN, DRWN, AB
FP.EX ----- FP for original and latest publication
FPALL ----- PI, TI, IN, INA, PA, PAA, PAT, PETRM, DCD, AI,
            RLI, PRAI, IC, ICM, ICS, INCL, INCLM, INCLS, NCL, NCLM,
            NCLS, EXF, REP, REN, ARTU, EXNAM, LREP, CLMN, DRWN, AB,
            PARN, SUMM, DRWD, DETD, CLM
FPBIB ----- PI, TI, IN, INA, PA, PAA, PAT, PTERM, DCD, AI,
            RLI, PRAI, REP, REN, EXNAM, LREP, CLM, CLMN, DRWN
FHITSTR ---- HIT RN, its text modification, its CA index name, and
            its structure diagram
FPG ----- FP plus PAGE.DRAW
GI. ----- PN and page image numbers
HIT ----- All fields containing hit terms
HITRN ----- HIT RN and its text modification
HITSTR ---- HIT RN, its text modification, its CA index name, and
            its structure diagram
IABS ----- ABS, indented with text labels
IALL ----- ALL, indented with text labels
IALLG ----- IALL plus PAGE.DRAW
IBIB ----- BIB, indented with text labels
IBIB.EX ---- IBIB for original and latest publication
IBIBG ----- IBIB plus PAGE.DRAW
IMAX ---- MAX, indented with text labels
IMAX.EX ---- IMAX for original and latest publication
IND ----- INCL, INCLM, INCLS, NCL, NCLM, NCLS, IC, ICM, ICS,
            EXF, ARTU, OS, CC, SX, ST, IT
```

ISTD ----- STD, indented with text labels

```
KWIC ----- All hit terms plus 20 words on either side
MAX ----- AN, TI, IN, INA, PA, PAA, PAT, PI, AI, PTERM, DCD,
             RLI, PRAI, DT, FS, REP, REN, EXNAM, LREP, CLMN, ECL,
             DRWN, AB, GOVI, PARN, SUMM, DRWD, DETD, CLM, INCL,
             INCLM, INCLS, NCL, NCLM, NCLS, IC, ICM, ICS,
             EXF, ARTU OS, CC, SX, ST, IT
MAX.EX ---- MAX for original and latest publication
OCC ----- List of display fields containing hit terms
SBIB ----- AN, TI, IN, INA, PA, PAA, PAT, PI, AI, RLI, PRAI,
             DT, FS, LN.CNT
SCAN ----- AN, TI, NCL, NCLM, NCLS, IC, ICM, ICS (random display
             without answer number. SCAN must be entered on the
             same line as DISPLAY, e.g., D SCAN)
STD ----- AN, TI, IN, INA, PA, PAA, PAT, PI, AI, RLI, PRAI,
             DT, FS, LN.CNT, INCL, INCLM, INCLS, NCL, NCLM, NCLS,
             IC, ICM, ICS, EXF (STD is the default)
STD.EX ---- STD for original and latest publication
TRIAL ---- AN, TI, INCL, INCLM, INCLS, NCL, NCLM, NCLS, IC,
             ICM, ICS
ENTER DISPLAY FORMAT (STD):STD
L7
     ANSWER 1 OF 26 USPATFULL
AN
       2003:123367 USPATFULL
       Method of treating metabolic disorders especially diabetes, or a disease
ΤI
       or condition associated with diabetes
       Gatlin, Marjorie Regan, Hoboken, NJ, United States
IN
       Ball, Michele Ann, Morris Plains, NJ, United States
       Mannion, Richard Owen, Mount Arlington, NJ, United States
       Karnachi, Anees Abdulquadar, Hillsborough, NJ, United States
       Guitard, Christiane, Hegenheim, FRANCE
       Allison, Malcolm, Basel, SWITZERLAND
       Novartis AG, Basel, SWITZERLAND (non-U.S. corporation)
PA ·
                               20030506
PΙ
       US 6559188
                          В1
       US 2000-663264
                               20000915 (9)
ΑI
                           20000407 (60)
PRAI
       US 2000-304196P
                           20000309 (60)
       US 2000-240918P
                           19990917 (60)
       US 1999-242911P
DT
       Utility
FS
       GRANTED
LN.CNT 2176
       INCLM: 514/641.000
INCL
       INCLS: 514/023.000; 514/057.000; 514/866.000
NCL
       NCLM:
              514/641.000
              514/023.000; 514/057.000; 514/866.000
       NCLS:
IC
       [7]
       ICM: A61K031-13
       ICS: A61K031-70; A61K031-715
       514/641; 514/866; 514/23; 514/57
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
```

=> d 17 1-26 bib, kwic

L7 ANSWER 1 OF 26 USPATFULL
AN 2003:123367 USPATFULL

TI Method of treating metabolic disorders especially diabetes, or a disease or condition associated with diabetes

IN Gatlin, Marjorie Regan, Hoboken, NJ, United States
Ball, Michele Ann, Morris Plains, NJ, United States
Mannion, Richard Owen, Mount Arlington, NJ, United States
Karnachi, Anees Abdulquadar, Hillsborough, NJ, United States

```
Guitard, Christiane, Hegenheim, FRANCE
       Allison, Malcolm, Basel, SWITZERLAND
       Novartis AG, Basel, SWITZERLAND (non-U.S. corporation)
PA
PΙ
       US 6559188
                          В1
                                20030506
       US 2000-663264
                                20000915 (9)
AΤ
       US 2000-304196P
                           20000407 (60)
PRAI
       US 2000-240918P
                           20000309 (60)
                           19990917 (60)
       US 1999-242911P
DT
       Utility
FS
       GRANTED
       Primary Examiner: Weddington, Kevin E.
EXNAM
LREP
       Thallemer, John D.
CLMN
       Number of Claims: 11
ECL
       Exemplary Claim: 1
DRWN
       0 Drawing Figure(s); 0 Drawing Page(s)
LN.CNT 2176
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
            . daily. Repaglinide is administered in a dosage of preferably
       0.01 to 8 mg per meal, more preferably about 0.2 to 5
       mg per meal, and most preferably 0.5 mg to 4
       mg per meal.
SUMM
       . . . dosage of englitazone or darglitazone is preferably in the
       range of about 0.05 to 50, more preferably about 0.05 to 5,
       mg/kg body weight of the patient per day, if the warm-blooded
       animal is a human. The dosage of AY-31637 is in. . . to 100, mg/kg
       body weight of the patient per day, if the warm-blooded animal is a
       human. The dosage of ciglitazone is in the range of about 0.25
       to 200, more preferably about 0.5 to 50, mg/kg body weight of the.
L7
     ANSWER 2 OF 26 USPATFULL
ΑN
       2001:226662 USPATFULL
TI
       Hypoglycemic thiazolidinediones and intermediates
       Clark, David A., East Lyme, CT, United States
TN
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
PΙ
       US 6329408
                          В1
                               20011211
ΑI
       US 1995-458071
                               19950601 (8)
       Division of Ser. No. US 1993-162027, filed on 1 Dec 1993, now abandoned
RLI
       Continuation of Ser. No. WO 1992-US5436, filed on 1 Jul 1992
       Continuation of Ser. No. US 1991-733771, filed on 22 Jul 1991, now
       abandoned
DT
       Utility
       GRANTED
FS
       Primary Examiner: McKane, Joseph K.; Assistant Examiner: D'Souza, Andrea
EXNAM
       Richardson, Peter C., Benson, Gregg C., Ronau, Robert T.
LREP
CLMN
       Number of Claims: 15
ECL
       Exemplary Claim: 1
       No Drawings
DRWN
LN.CNT 776
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
     ANSWER 3 OF 26 USPATFULL
T.7
       2001:158319 USPATFULL
ΑN
       Treating cancers associated with overexpression of class I family of
TI
       receptor tyrosine kinases
       Dannenberg, Andrew J., 7 Gracie Sq., Apt. 14A, New York, NY, United
IN
```

```
States 10028
       Subbaramaiah, Kotha, 43-23 Colden St., Apt. 17K, Flushing, NY, United
       States 11355
       US 6291496
                          В1
                               20010918
ΡI
                               19991227 (9)
ΑI
       US 1999-472179
DT
       Utility
FS
       GRANTED
      Primary Examiner: Eyler, Yvonne; Assistant Examiner: Andres, J.
EXNAM
       Number of Claims: 28
CLMN
       Exemplary Claim: 1
ECL
DRWN
       2 Drawing Figure(s); 2 Drawing Page(s)
LN.CNT 796
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      A patient with HER-2/neu positive breast cancer is treated with
       ciglitazone or troglitazone or pioglitazone or rosiglitazone for
       adjuvant therapy at an oral dose of 5 mg/kg twice
       per day for five years after a mastectomy. Recurrence of breast cancer
       does not occur.
       When GW 7845 or the compound of structure (II) or (III) is given at an
DETD
       oral dose of 5 mg/kg twice per day for five years
       after a mastectomy in place of the thiazolidinedione, recurrence of
       breast cancer does not.
       . . . mg intravenous HERCEPTIN.RTM., then 10 weekly doses of 125 mg
DETD
       each IV. The patient also received an oral dose of 5
       mg/kg ciglitazone or troglitazone or pioglitazone or
       rosiglitazone twice daily for one year or 5 mg/kg of
       GW 7845 or compound of structure (II) or (III) twice daily for one year.
       Recurrence of breast cancer does.
            . a mastectomy is performed on a patient with HER-2/neu positive
DETD
       breast cancer. The patient is treated with oral doses of 5
       mg/kg of ciglitazone or troglitazone or pioglitazone
       or rosiglitazone twice daily or \mathbf{5} \mathbf{mg}/\mathrm{kg} of GW 7845
       or compound of structure (II) or (III) twice daily. A reduced tumor
       burden is noted.
       After failure of prior chemotherapy regimens, the patient is treated
DETD
       with an oral dose of 5 mg/kg of ciglitazone
       or troglitazone or pioglitazone or rosiglitazone twice daily or
       5 mg/kg of GW 7845 or compound of structure (II) or
       (III) twice daily and HERCEPTIN.RTM. at a loading dose of 250.
            . to the bowel wall; there is no evidence of extracolonic cancer.
DETD
       The patient is treated with an oral dose of 5 mg/kg
       of ciglitazone or troglitazone or pioglitazone or
       rosiglitazone twice daily or 5 mg/kg of GW 7845 or
       compound of the structure (II) or (III) twice daily, for five years.
       Recurrence of colon cancer:
DETD
            . non-small cell lung carcinoma metastasized to liver, where EGFR
       is determined to be overexpressed, is treated with oral doses of
       5 mg/kg of ciglitazone or troglitazone or
       pioglitazone or rosiglitazone twice daily or 5 mg/kg
       of GW 7845 or compound of structure (II) or (III) twice daily. A reduced
       tumor burden is noted. After one.
       . . . 60 mg/m.sup.2. A reduced tumor burden is noted. At the end of
DETD
       the 12 week period, treatment is started with 5 mg
       /kg twice daily by oral administration of ciglitazone or
       troglitazone or pioglitazone or rosiglitazone or GW 7845 or compound of
       structure (II) or (E) for 12 weeks. A.
       . . . a dose of 175 mg/m.sup.2 administration intravenously every
DETD
       three weeks in further combination regimen with oral doses twice daily
       of 5 mg/kg of ciglitazone or troglitazone
       or pioglitazone or rosiglitazone or GW 7845 or compound of structure (I)
       or (III). A reduced tumor burden.
DETD
       A patient with EGFR positive breast cancer is treated with
```

ciglitazone or troglitazone or pioglitazone or rosiglitazone or
GW 7845 or compound of structure (II) or (III) at an oral dose of
5 mg/kg twice per daily for five years after a
mastectomy. Recurrence of breast cancer dose not occur.

```
ANSWER 4 OF 26 USPATFULL
L7
AN
       1998:69051 USPATFULL
       Secondary amines as antidiabetic and antiobesity agents
TI
       Dow, Robert L., Waterford, CT, United States
IN
       Wright, Stephen W., Old Lyme, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
                               19980616
       US 5767133
PΙ
       WO 9429290 19941222
       US 1995-569152
                               19951214 (8)
ΑI
       WO 1994-IB117
                               19940520
                               19951214
                                          PCT 371 date
                               19951214 PCT 102(e) date
DT
       Utility
       Granted
FS
       Primary Examiner: Ivy, C. Warren; Assistant Examiner: Aulakh, Charandit
EXNAM
       Richardson, Peter C., Benson, Gregg C., Ronau, Robert T.
LREP
       Number of Claims: 52
CLMN
ECL
       Exemplary Claim: 1
       No Drawings
DRWN
LN.CNT 2986
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . is collected via an ocular bleed prior to any treatment. The
SUMM
       blood sample is immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals are then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs are.
     ANSWER 5 OF 26 USPATFULL
L7
AN
       1998:65249 USPATFULL
       Hypoglycemic hydroxyurea derivatives
TΙ
       Goldstein, Steven Wayne, Mystic, CT, United States
IN
       McDermott, Ruth Elsbree, Salem, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PΆ
                               19980609
PΙ
       US 5763467
ΑI
       US 1997-840179
                               19970411 (8)
       Division of Ser. No. US 1995-544010, filed on 10 Oct 1995, now patented,
       Pat. No. US 5646168 which is a division of Ser. No. US 1995-391308,
       filed on 17 Feb 1995, now patented, Pat. No. US 5463070 which is a
       continuation of Ser. No. US 1994-279322, filed on 22 Jul 1994, now
       abandoned which is a division of Ser. No. US 1993-983538, filed on 22
       Feb 1993, now patented, Pat. No. US 5334604 which is a continuation of
      Ser. No. US 1990-572745, filed on 23 Aug 1990, now abandoned
DT
       Utility
       Granted
       Primary Examiner: Daus, Donald G.
EXNAM
       Richardson, Peter C., Benson, Gregg C., Jones, James T.
       Number of Claims: 15
CLMN
       Exemplary Claim: 1
ECL
       No Drawings
DRWN
LN.CNT 1122
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
SUMM
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
```

held on ice for metabolite analysis. Animals were then dosed daily for five days with drug (5-50 mg/kg), a positive control (50 mg/kg) of ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm. Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were. ANSWER 6 OF 26 USPATFULL 97:59226 USPATFULL Hypoglycemic hydroxyurea derivatives Goldstein, Steven Wayne, 176 Bel-Aire Dr., Mystic, CT, United States McDermott, Ruth Elsbree, 38 Meadow La., Salem, CT, United States 19970708 US 5646168 US 1995-544010 19951010 (8) Division of Ser. No. US 1995-391308, filed on 17 Feb 1995, now patented, Pat. No. US 5463070 which is a continuation of Ser. No. US 1994-279322, filed on 22 Jul 1994, now abandoned which is a division of Ser. No. US 1993-983538, filed on 22 Feb 1993, now patented, Pat. No. US 5334604 which is a continuation of Ser. No. US 1990-572745, filed on 23 Aug 1990, now abandoned Utility Granted Primary Examiner: Daus, Donald G. Number of Claims: 6 Exemplary Claim: 1 No Drawings LN.CNT 1076 CAS INDEXING IS AVAILABLE FOR THIS PATENT was collected via an ocular bleed prior to any treatment. The blood sample was immediately diluted 1:5 with saline containing 2. 5 mg/ml sodium fluoride and 2% sodium heparin, and held on ice for metabolite analysis. Animals were then dosed daily for five days with drug (5-50 mg/kg), a positive control (50 mg/kg) of ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm. Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were. ANSWER 7 OF 26 USPATFULL 96:29574 USPATFULL Thiazolidinedione compounds Regnier, Gilbert, Chatenay Malabry, France Charton, Yves, Sceaux, France Duhault, Jacques, Croissy Sur Seine, France Espinal, Joseph, Levallois Perret, France ADIR et Compagnie, Courbevoie, France (non-U.S. corporation) US 5506245 19960409 US 1995-374970 19950119 (8) Continuation-in-part of Ser. No. US 1993-133898, filed on 12 Oct 1993, now abandoned FR 1992-12123 19921012 Utility Granted Primary Examiner: Gerstl, Robert Hueschen, Gordon W. Number of Claims: 4 Exemplary Claim: 1 No Drawings

Doses causing the same hypoglycaemic effect in ob/ob mice

TABLE 1

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Dose mg/kg/day

L7

ANTI

IN

PΙ

ΑI

DT FS

EXNAM CLMN

ECL

DRWN

DETD

L7

ANΤI

IN

PA PΙ

ΑI

RLI

PRAI DT

EXNAM LREP

CLMN

ECL

DRWN

DETD

LN.CNT 446

FS

RLI

```
Feb 1993, now patented, Pat. No. US 5334604 which is a continuous Ser. No. US 1990-572745, filed on 23 Aug 1990, now abandoned Utility
FS Granted
EXNAM Primary Examiner: Daus, Donald G.
LREP Richardson, Peter C., Benson, Gregg C., Jones, James T.
CLMN Number of Claims: 5
ECL Exemplary Claim: 1
```

```
DRWN
       No Drawings
LN.CNT 1076
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
            . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
L7
     ANSWER 10 OF 26 USPATFULL
AN
       95:69300 USPATFULL
       3-aryl-2-hydroxypropionic acid derivatives and analogs as hypoglycemic
TΙ
IN
       Hulin, Bernard, Essex, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
PΙ
       US 5438074
                                19950801
       US 1993-163781
                                19931206 (8)
ΑI
       Division of Ser. No. US 1992-980404, filed on 24 Nov 1992, now patented,
RLI
       Pat. No. US 5306726 which is a continuation-in-part of Ser. No. US
       1990-537673, filed on 14 Jun 1990, now patented, Pat. No. US 5089514
DT
       Utility
FS
       Granted
       Primary Examiner: Chan, Nicky
EXNAM
       Richardson, Peter C., Benson, Gregg C., Ronau, Robert T.
LREP
CLMN
       Number of Claims: 8
ECL
       Exemplary Claim: 1,8
DRWN
       No Drawings
LN.CNT 1706
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
L7
     ANSWER 11 OF 26 USPATFULL
ΑN
       95:27328 USPATFULL
TI
       Thiazolidinedione hypoglycemic agents
       Goldstein, Steven W., Mystic, CT, United States
IN
       Hulin, Bernard, Essex, CT, United States
       Pfizer, Inc., New York, NY, United States (U.S. corporation)
PA
PΙ
       US 5401761
                                19950328
ΑI
       US 1993-162075
                                19931209 (8)
DT
       Utility
FS
       Granted
EXNAM
       Primary Examiner: Gerstl, Robert
LREP
       Richardson, Peter C., Benson, Gregg C., Brokke, Mervin E.
CLMN
       Number of Claims: 14
       Exemplary Claim: 1
ECL
DRWN
       No Drawings
LN.CNT 643
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
```

ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.

```
Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
L7
     ANSWER 12 OF 26 USPATFULL
       94:66495 USPATFULL
AN
TΙ
       Hypoglycemic hydroxyurea derivatives
       Goldstein, Steven W., Mystic, CT, United States
ΤN
       McDermott, Ruth E., Salem, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
PΙ
       US 5334604
                                19940802
       US 1993-983538
                                19930222
ΑI
       WO 1991-US4352
                                19910626
                                19930222
                                          PCT 371 date
                                19930222
                                          PCT 102(e) date
       Continuation of Ser. No. US 1990-572745, filed on 23 Aug 1990, now
RLI
       abandoned
DT
       Utility
FS
       Granted
EXNAM
       Primary Examiner: Daus, Donald G.
       Richardson, Peter C., Benson, Gregg C., McFarlin, D. Stuart
LREP
       Number of Claims: 6
CLMN
ECL
       Exemplary Claim: 1
       No Drawings
DRWN
LN.CNT 1045
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
DETD
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 mg/kg), a positive control (50 mg/kg) of ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
     ANSWER 13 OF 26 USPATFULL
T.7
       94:62459 USPATFULL
AN
       Thiazolidinedione derivatives as hypoglycemic agents
TI
       Clark, David A., East Lyme, CT, United States
IN
       Goldstein, Steven W., Mystic, CT, United States
       Hulin, Bernard, Essex, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PΑ
PΙ
       US 5330998
                                19940719
ΑI
       US 1990-566437
                                19900814 (7)
       WO 1988-US745
                                19880308
                                19900814
                                           PCT 371 date
                                19900814
                                          PCT 102(e) date
DT
       Utility
FS
       Granted
EXNAM
       Primary Examiner: Gerstl, Robert
       Richardson, Peter C., Benson, Gregg C., Brokke, Mervin E.
LREP
       Number of Claims: 28
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 1144
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
SUMM
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
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L7

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AN
       94:35602 USPATFULL
       3-aryl-2-hydroxypropionic acid derivatives and analogs as hypoglycemic
TΙ
IN
       Hulin, Bernard, Essex, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PΑ
                               19940426
PΙ
       US 5306726
       US 1992-980404
                               19921124 (7)
ΑI
       which is a continuation-in-part of Ser. No. US 1990-537673, filed on 14
RLI
       Jun 1990, now patented, Pat. No. US 5089514
DΤ
       Utility
FS
       Granted
EXNAM
       Primary Examiner: Daus, Donald G.
LREP
       Richardson, Peter C., Benson, Gregg C., Ronau, Robert T.
CLMN
       Number of Claims: 4
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 1686
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
     ANSWER 15 OF 26 USPATFULL
ь7
       93:52600 USPATFULL
ΑN
       Thiazolidinedione hypoglycemic agents
TI
       Clark, David A., East Lyme, CT, United States
IN
       Goldstein, Steven W., Mystic, CT, United States
       Holland, Gerald F., Old Lyme, CT, United States
       Hulin, Bernard, Essex, CT, United States
       Rizzi, James P., Waterford, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
PΙ
       US 5223522
                               19930629
ΑI
       US 1992-857039
                               19920324 (7)
       Division of Ser. No. US 1991-679898, filed on 3 Apr 1991, now patented,
RLT
       Pat. No. US 5120754 which is a division of Ser. No. US 1990-566436,
       filed on 14 Aug 1990, now patented, Pat. No. US 5061717
DΤ
       Utility
FS
       Granted
       Primary Examiner: Rotman, Alan L.
EXNAM
       Richardson, Peter C., Benson, Gregg C., Brokke, Mervin E.
LREP
CLMN
       Number of Claims: 6
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 1598
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
     ANSWER 16 OF 26 USPATFULL
L7
ΑN
       92:57744 USPATFULL
       Hypoglycemic thiazolidinedione derivatives
ΤI
     . Clark, David A., Groton, CT, United States
ΙN
       Goldstein, Steven W., Groton, CT, United States
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Hulin, Bernard, Groton, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
       US 5130379
                               19920714
PΙ
       US 1991-674833
                               19910326 (7)
ΑI
       Division of Ser. No. US 1990-477261, filed on 8 Feb 1990, now patented,
RLI
       Pat. No. US 5036079 which is a continuation-in-part of Ser. No. US
       1989-438490, filed on 7 Dec 1989, now abandoned
DT
       Utility
FS
       Granted
      Primary Examiner: Rotman, Alan L.
EXNAM
       Richardson, Peter C., Lumb, J. Trevor, Brokke, Mervin E.
LREP
       Number of Claims: 37
CLMN
       Exemplary Claim: 1
ECL
DRWN ·
       No Drawings
LN.CNT 1362
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
SUMM
       . . . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50~\text{mg/kg}), a positive control (50~\text{mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
     ANSWER 17 OF 26 USPATFULL
L7
AN
       92:47080 USPATFULL
TΙ
       Thiazolidinedione hypoglycemic agents
       Clark, David A., East Lyme, CT, United States
IN
       Goldstein, Steven W., Mystic, CT, United States
       Holland, Gerald F., Old Lyme, CT, United States
       Hulin, Bernard, Essex, CT, United States
       Rizzi, James P., Waterford, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
       US 5120754
                                19920609
PΙ
ΑI
       US 1991-679898
                                19910403 (7)
       Division of Ser. No. US 1990-566436, filed on 14 Aug 1990, now patented,
RLI
       Pat. No. US 5061717
DT
       Utility
       Granted
FS
       Primary Examiner: Rotman, Alan L.
EXNAM
       Richardson, Peter C., Lumb, J. Trevor, Brokke, Mervin E.
LREP
       Number of Claims: 22
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 1604
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
            . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
     ANSWER 18 OF 26 USPATFULL
L7
       92:12971 USPATFULL
AN
       3-coxazolyl [phenyl, chromanyl or benzofuranyl]-2-hydroxypropionic acid
ΤI
       derivatives and analogs as hypoglycemic agents
       Hulin, Bernard, Groton, CT, United States
ΙN
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
                                19920218
PΙ
       US 5089514
                                19900614 (7)
       US 1990-537673
ΑI
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DT
       Utility
       Granted
FS
       Primary Examiner: Daus, Donald G.
EXNAM
       Richardson, Peter C., Lumb, J. Trevor, McFarlin, D. Stuart
       Number of Claims: 42
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 1169
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
L7
     ANSWER 19 OF 26 USPATFULL
ΑN
       91:89062 USPATFULL
       Thiazolidinedione hypoglycemic agents
TI ·
       Clark, David A., East Lyme, CT, United States
TN
       Goldstein, Steven W., Mystic, CT, United States
       Holland, Gerald F., Old Lyme, CT, United States
       Hulin, Bernard, Essex, CT, United States
       Rizzi, James P., Waterford, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
                                19911029
PΙ
       US 5061717
                                19900814 (7)
ΑI
       US 1990-566436
       WO 1988-US744
                                19880308
                                19900814
                                          PCT 371 date
                                19900814 PCT 102(e) date
       Utility
DT
FS
       Granted
EXNAM
       Primary Examiner: Rotman, Alan L.
LREP
       Richardson, Peter C., Lumb, J. Trevor, Brokke, Mervin E.
       Number of Claims: 13
CLMN
       Exemplary Claim: 1,11
ECL
DRWN
       No Drawings
LN.CNT 1594
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
             . was collected via an ocular bleed prior to any treatment. The
SUMM
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
L7
     ANSWER 20 OF 26 USPATFULL
ΑN
       91:62804 USPATFULL
       Oxa- and thiazolidinedione hypoglycemic and hypocholesterolemic agents
ΤI
       Goldstein, Steven W., Mystic, CT, United States
IN
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
                                19910806
PΙ
       US 5037842
       US 1990-533615
                                19900605 (7)
ΑI
       Utility
DT
       Granted
       Primary Examiner: Daus, Donald G.
EXNAM
LREP
       Richardson, Peter C., Lumb, J. Trevor, McManus, James M.
CLMN
       Number of Claims: 9
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
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LN.CNT 586
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
            . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days drug (5-50 mg/kg), a positive control (50 mg/kg) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull , vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs.
L7
    ANSWER 21 OF 26 USPATFULL
AN
       91:60822 USPATFULL
       Hypoglycemic thiazolidinedione derivatives
TI
       Clark, David A., East Lyme, CT, United States
IN
       Goldstein, Steven W., Mystic, CT, United States
       Hulin, Bernard, Essex, CT, United States
PA
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PΙ
       US 5036079
                               19910730
                               19900208 (7)
ΑI
       US 1990-477261
       Continuation-in-part of Ser. No. US 1989-438490, filed on 7 Dec 1989,
RLI
       now abandoned
DT
       Utility
FS
       Granted
       Primary Examiner: Rotman, Alan L.
EXNAM
       Richardson: Peter C., Lumb, J. Trevor, Blackwood, Robert K.
LREP
       Number of Claims: 24
CLMN
ECL
       Exemplary Claim: 1,21
DRWN
       No Drawings
LN.CNT 1330
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
SUMM
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 mg/kg), a positive control (50 mg/kg) of
       ciglitazone; U.S. Pat. No. 4,467,902; Sohda et al., Chem. Pharm.
       Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
    ANSWER 22 OF 26 USPATFULL
L7
ΑN
       90:85644 USPATFULL
TI
       Oxazolidin-2-one derivatives as hypoglycemic agents
       Clark, David A., East Lyme, CT, United States
IN
       Johnson, Michael R., Chapel Hill, NC, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
                               19901106
       US 4968707
PΙ
       WO 8809661 19881215
ΑI
       US 1989-460848
                               19891115 (7)
       WO 1987-US1356
                               19870610
                               19891115
                                         PCT 371 date
                               19891115
                                        PCT 102(e) date
DT
       Utility
FS
       Granted
     Primary Examiner: Daus, Donald G.
EXNAM
       Richardson, Peter C., Lumb, J. Trevor, Blackwood, Robert K.
LREP
       Number of Claims: 22
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 913
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
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Bull., vol. 32, pp. 4460-4465, 1984), or vehicle. All drugs were.
L7
     ANSWER 23 OF 26 USPATFULL
AN
       90:6005 USPATFULL
       Novel naphthalenyl-3H-1,2,3,5-oxathiadiazole 2-oxides useful as
ΤI
       antihyperglycemic agents
       Lombardo, Louis J., South Plainfield, NJ, United States
IN
       Alessi, Thomas R., Monmouth Junction, NJ, United States
       American Home Products Corporation, New York, NY, United States (U.S.
PA
       corporation)
                               19900123
PΙ
       US 4895861
ΑI
       US 1989-341609
                               19890421 (7)
DT
       Utility
FS
       Granted
       Primary Examiner: Gerstl, Robert
EXNAM
       Patton, Walter
LREP
       Number of Claims: 16
CLMN
       Exemplary Claim: 1
ECL
DRWN
       No. Drawings
LN.CNT 867
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . lower blood glucose levels in diabetic mice. For example,
       4-[1-(5-bromo-2-naphthalenyl)ethyl]-3H-1,2,3,5-oxathiadiazole 2-oxide,
       the compound of Example 1, at a dose of 5 mg/kg/day
       give comparable results to ciglitazone at 100 mg/kg/day.
                                             . . 20
                                                                 137-138 (dec.)
DETD
       2-S0.sub.2 CH.sub.2
Η
                 20
                                170-171
       2-OCH.sub.2
Η
                 20
                       4
                               101-103
                               146-147 (dec.)
    Η
       2-CHCH
                 5
                       4
  Ciglitazone
                   100
                         -33
     ANSWER 24 OF 26 USPATFULL
L7
       88:52199 USPATFULL
AN
       N-(1H-tetrazol-5-yl-alkylphenyl)polyfluoroalkanamides
ΤI
       Kees, Kenneth L., West Chester, PA, United States
IN
       American Home Products Corporation, New York, NY, United States (U.S.
PA
       corporation)
                               19880816
PΙ
       US 4764623
                               19870615 (7)
       US 1987-62270
ΑI
DT
       Utility
FS
       Granted
       Primary Examiner: Hollrah, Glennon H.; Assistant Examiner: Springer, D.
EXNAM
LREP
       Jackson, Richard K.
CLMN
       Number of Claims: 17
       Exemplary Claim: 1
ECL
DRWN
       No Drawings
LN.CNT 701
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . to be administered to a mammal suffering from excessive blood
DETD
       levels of glucose and/or insulin in an amount from about 5
       mg./kg. to about 300 mg./kg. body weight or more per day. An
       optimum dosing regimen to achieve the desired therapeutic response.
          problem of precipitating hypoglycemic shock. In addition, the
       compounds of Examples 1 and 3 are more effective than the standard
       ciglitazone in reducing blood glucose levels.
```

held on ice for metabolite analysis. Animals were then dosed daily for five days with drug (5-50 mg/kg), a positive control (50 mg/kg of ciglitazone; U.S. Pat. No. 4,461,902; Sohda et al., Chem. Pharm.

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AN
       88:24416 USPATFULL
       Hypoglycemic thiazolidinediones
ΤI
       Eggler, James F., Stonington, CT, United States
IN
       Holland, Gerald F., Old Lyme, CT, United States
       Johnson, Michael R., Gales Ferry, CT, United States
       Volkmann, Robert A., Ledyard, CT, United States
       Pfizer Inc., New York, NY, United States (U.S. corporation)
PA
       US 4738972
                                19880419
PΙ
ΑI
       US 1987-67899
                                19870626 (7)
       Continuation-in-part of Ser. No. US 1986-10081, filed on 29 Dec 1986,
RLI
       now patented, Pat. No. US 4703052
DT
       Utility
FS
       Granted
       Primary Examiner: Gerstl, Robert
EXNAM
       Richardson, Peter C., Frost, Albert E., Blackwood, Robert K.
LREP
CLMN
       Number of Claims: 23
       Exemplary Claim: 1
ECL
DRWN
       No Drawings
LN.CNT 1318
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . was collected via an ocular bleed prior to any treatment. The
SUMM
       blood sample was immediately diluted 1:5 with saline containing 2.
       5 mg/ml sodium fluoride and 2% sodium heparin, and
       held on ice for metabolite analysis. Animals were then dosed daily for
       five days with drug (5-50 \text{ mg/kg}), a positive control (50 \text{ mg/kg}) of
       ciglitazone; U.S. Pat. No. 4.467,,902; Sohda et al., Chem.
       Pharm. Bull., vol. 32, pp. 4460-4465, 1984), or vehicle All drugs were.
     ANSWER 26 OF 26 USPATFULL
L7
       88:13313 USPATFULL
AN
       Hypoglycemic thiazolidinediones
TI
       Kees, Kenneth L., West Chester, PA, United States
IN
       Cheeseman, Robert S., Swedeland, PA, United States
       American Home Products Corporation, New York, NY, United States (U.S.
PA
       corporation)
       US 4728739
                                19880301
PΙ
ΑI
       US 1987-62268
                                19870615 (7)
DT
       Utility
FS
       Granted
       Primary Examiner: Gerstl, Robert
EXNAM
       Jackson, Richard K.
LREP
       Number of Claims: 4
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 181
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
DETD
                      TABLE
                 Dose
Experi-
ment Drug
                 (mg/kg)
                          Glucose (mg/dl)
                                     Insulin (.mu.ml)
                          154 ..+-.
1
      control
                                       182 + 9
                                 10
        Ciglitazone
                 75
                                7*
                                       57 .+-.
      Example 1 75
                          94 .+-.
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ANSWER 25 OF 26 USPATFULL

L7

				3*	45 .+ 2*
2	control		139 .+-	8	167 .+ 14
	Ciglita	.zone 5	127 .+-	 5	156 .+
	Example 1	. 5	105 .+	 6*	12 63 .+ 7*
3	control		137 .+-	 7	168 .+ 24
					24
	Ciglita	5	101 .+		170 .+
		20	97 .+-		116 .+
		75	92 .+-	• 4*.	59 .+
	+				
				3*	51 .+
		75	87 .+-	3*	31 .+
4	control		120 .+	 7	229 . +
	Ciglita	7000			
	CIGILLA	20	76 .+-	_	•
			76 .+-		183 .+ 20
	Example 2	20	81 .+-	· 7*	137 .+ 16*

^{*}Significantly different from control. . .

DETD . . . to be administered to a mammal suffering from excessive blood levels of glucose and/or insulin in an amount from about 5 mg/kg to about 300 mg/kg body weight or more per day. An optimum dosing regimen to achieve the desired therapeutic response. . .